AMENDMENTS TO THE CLAIMS

1. (Currently Amended) A gel matrix comprising a hydrated an electrophoresis gel

comprising pores having a size to sieve molecules of a desired size range by electrophoresis expressions and the size range by electrophoresis expressions.

magnetophoresis—and one or more SERS-enhancing nanoparticles stationary within the

electrophoresis gel,

wherein the gel matrix has a thickness sufficient to perform electrophoresis-the-nanoparticles

are composite organicinorganic nanoparticle (COINS) comprising a core and a surface, wherein the

core comprises a metallic colloid comprising a first metal and a Raman active organic/compound.

2. (Original) The gel matrix of claim 1 comprising a plurality of the nanoparticles to

provide a plurality of unique optical signatures.

3. (Original) The gel matrix of claim 2, wherein the SERS-enhancing nanoparticles

comprise one or more Raman-active tags independently selected from the group consisting of

nucleic acids, nucleotides, nucleotide analogs, base analogs, fluorescent dyes, peptides, amino acids,

modified amino acids, organic moieties, quantum dots, carbon nanotubes, fullerenes, metal

nanoparticles, electron dense particles and crystalline particles.

4. (Original) The gel matrix of claim 1, wherein at least one of the nanoparticles has a net

charge.

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5. (Original) The gel matrix of claim 1, wherein the nanoparticles each provide a unique

SERS-signal that is correlated with binding specificity of the probe of the nanoparticle.

6. (Original) The gel matrix of claim 1, wherein the Raman-active tag comprises adenine or

an analog thereof.

7-9. (Canceled)

10. (Original) The gel matrix of claim 1, wherein the probe is selected from antibodies,

antigens, polynucleotides, oligonucleotides, receptors and ligands.

11. (Original) The gel matrix of claim 10, wherein the probe comprises a polynucleotide.

12. (Previously presented) The gel matrix of claim 1, wherein any of the nanoparticles may

further comprise a fluorescent label that contributes to the optical signature.

13-32. (Canceled)

33. (Currently Amended) A system for detecting an analyte in a sample comprising a gel

matrix comprising a hydrated an electrophoresis gel comprising pores having a size to sieve

molecules of a desired size range by electrophoresis or magnetophoresis and one or more SERS-

enhancing nanoparticles stationary within the <u>electrophoresis</u> gel, the SERS-enhancing

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nanoparticles within the electrophoresis gel having an attached probe that binds specifically to an

analyte; a sample containing at least one analyte; and

an optical detection system suitable for detecting SERS signals from the nanoparticles,

wherein the gel matrix is thick enough to perform electrophoresis the nanoparticles are composite

organicinorganic nanoparticle (COINS) comprising a core and a surface, wherein the core

comprises a metallic colloid comprising a first metal and a Raman active organic/compound.

34. (Original) The system of claim 33, further comprising a computer comprising an

algorithm for analysis of the SERS signals obtained from the sample.

35-93. (Canceled)

94. (Currently amended) The gel matrix of claim 1, wherein the SERS-enhancing

nanoparticles within the electrophoresis gel have an attached probe that binds specifically to an

analyte.

95. (New) A gel matrix comprising a magnetophoresis gel comprising pores having a size to

sieve molecules of a desired size range by magnetophoresis and one or more SERS-enhancing

nanoparticles stationary within the magnetophoresis gel,

wherein the gel matrix has a thickness sufficient to perform electrophoresis.

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96. (New) A system for detecting an analyte in a sample comprising a gel matrix comprising

a magnetophoresis gel comprising pores having a size to sieve molecules of a desired size range by

electrophoresis or magnetophoresis and one or more SERS-enhancing nanoparticles stationary

within the magnetophoresis magnetophoresis gel, the SERS-enhancing nanoparticles within the

magnetophoresis gel having an attached probe that binds specifically to an analyte; a sample

containing at least one analyte; and

an optical detection system suitable for detecting SERS signals from the nanoparticles,

wherein the gel matrix is thick enough to perform magnetophoresis.

97. (New) The gel matrix of claim 33 comprising a plurality of the nanoparticles to

provide a plurality of unique optical signatures.

98. (New) The gel matrix of claim 97, wherein the SERS-enhancing nanoparticles

comprise one or more Raman-active tags independently selected from the group consisting of

nucleic acids, nucleotides, nucleotide analogs, base analogs, fluorescent dyes, peptides, amino acids,

modified amino acids, organic moieties, quantum dots, carbon nanotubes, fullerenes, metal

nanoparticles, electron dense particles and crystalline particles.

99. (New) The gel matrix of claim 33, wherein at least one of the nanoparticles has a net

charge.

100. (New) The gel matrix of claim 33, wherein the nanoparticles each provide a unique

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SERS-signal that is correlated with binding specificity of the probe of the nanoparticle.

101. (New) The gel matrix of claim 33, wherein the Raman-active tag comprises adenine or

an analog thereof.

102. (New) The gel matrix of claim 33, wherein the probe is selected from antibodies,

antigens, polynucleotides, oligonucleotides, receptors and ligands.

103. (New) The gel matrix of claim 102, wherein the probe comprises a polynucleotide.

104. (New) The gel matrix of claim 33, wherein any of the nanoparticles may

further comprise a fluorescent label that contributes to the optical signature.

105. (New) The gel matrix of claim 95 comprising a plurality of the nanoparticles to

provide a plurality of unique optical signatures.

106 (New) The gel matrix of claim 105, wherein the SERS-enhancing nanoparticles

comprise one or more Raman-active tags independently selected from the group consisting of

nucleic acids, nucleotides, nucleotide analogs, base analogs, fluorescent dyes, peptides, amino acids,

modified amino acids, organic moieties, quantum dots, carbon nanotubes, fullerenes, metal

nanoparticles, electron dense particles and crystalline particles.

107. (New) The gel matrix of claim 95, wherein at least one of the nanoparticles has a net

charge.

108. (New) The gel matrix of claim 95, wherein the nanoparticles each provide a unique

SERS-signal that is correlated with binding specificity of the probe of the nanoparticle.

109. (New) The gel matrix of claim 95, wherein the Raman-active tag comprises adenine or

an analog thereof.

110. (New) The gel matrix of claim 95, wherein the probe is selected from antibodies,

antigens, polynucleotides, oligonucleotides, receptors and ligands.

111. (New) The gel matrix of claim 110, wherein the probe comprises a polynucleotide.

112. (New) The gel matrix of claim 95, wherein any of the nanoparticles may

further comprise a fluorescent label that contributes to the optical signature.

113. (New) The gel matrix of claim 96 comprising a plurality of the nanoparticles to

provide a plurality of unique optical signatures.

114. (New) The gel matrix of claim 113, wherein the SERS-enhancing nanoparticles

comprise one or more Raman-active tags independently selected from the group consisting of

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nucleic acids, nucleotides, nucleotide analogs, base analogs, fluorescent dyes, peptides, amino acids,

modified amino acids, organic moieties, quantum dots, carbon nanotubes, fullerenes, metal

nanoparticles, electron dense particles and crystalline particles.

115. (New) The gel matrix of claim 96, wherein at least one of the nanoparticles has a net

charge.

116. (New) The gel matrix of claim 96, wherein the nanoparticles each provide a unique

SERS-signal that is correlated with binding specificity of the probe of the nanoparticle.

117. (New) The gel matrix of claim 96, wherein the Raman-active tag comprises adenine or

an analog thereof.

118. (New) The gel matrix of claim 96, wherein the probe is selected from antibodies,

antigens, polynucleotides, oligonucleotides, receptors and ligands.

119. (New) The gel matrix of claim 118, wherein the probe comprises a polynucleotide.

120. (New) The gel matrix of claim 96, wherein any of the nanoparticles may

further comprise a fluorescent label that contributes to the optical signature.